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NUTRITIONAL INTERVENTIONS FOR CANCER-INDUCED CACHEXIA





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Abu Dhabi, United Arab of Emirates 03 - 06 March 2016



Clinical Guidelines



Guidelines for the Provision and Assessment of Nutrition Support Therapy in the Adult Critically Ill Patient: Society of Critical Care Medicine (SCCM) and American Society for Parenteral and Enteral Nutrition (A.S.P.E.N.) Journal of Parenteral and Enteral Nutrition Volume 40 Number 2 February 2016 159–211 © 2016 American Society for Parenteral and Enteral Nutrition and Society of Critical Care Medicine DOI: 10.1177/0148607115621863 jpen.sagepub.com hosted at online.sagepub.com

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\$SAGE

Preferred route of feeding

Healthy meal

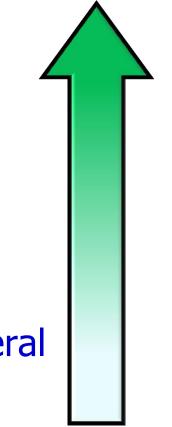
Oral

Enteral gastric

Enteral duodenal/jejunal

Small amount Enteral rest Parenteral

Total parenteral



Overview



- Weight loss to cancer-induced cachexia
- Specific nutrition for cancer patients -Nutrition in Cancer Care
- Summary

Overview

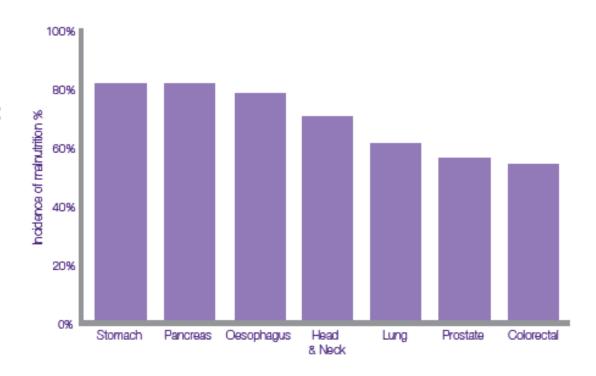


- Weight loss to cancer-induced cachexia
- Specific nutrition for cancer patients -Nutrition in Cancer Care
- Summary

Prevalence of Weight loss in Cancer



- 30-85% dependent on:
- Tumor (type & stage)
- Treatment
- Age
- Individual susceptibility
- Method of assessment



During the course of the disease there is weight loss of >10% in up to 45% of patients





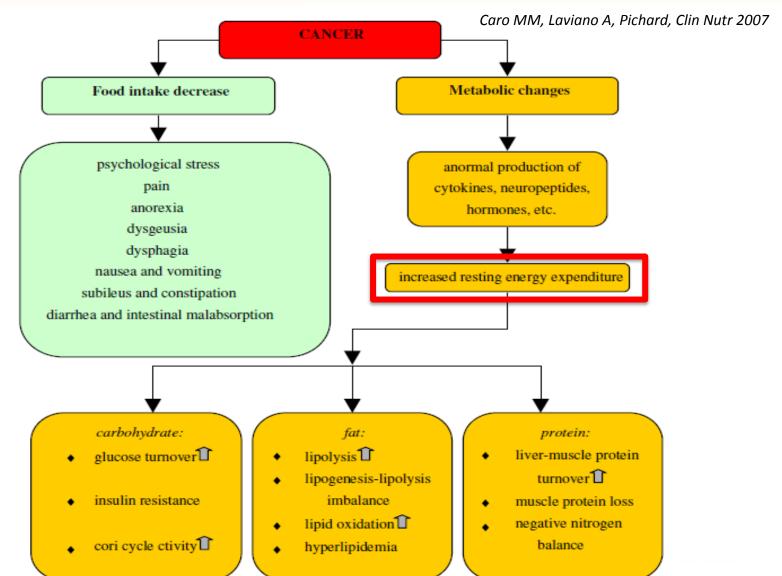
30-85% dependent on:
 Tumor (type & stage)
 Treatment
 Age
 Individual susceptibility
 Method of assessment

Over 20 % of death are due simply to malnutrition and host tissue wasting 50% of newly diagnosed cancer patients are anorexic

Nutrition; 12: 358-371, 1996

Weight loss in cancer patients







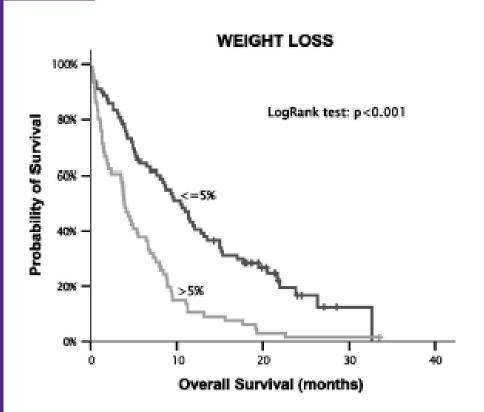
The higher rate of weight loss: the less survival



original article

Annals of Oncology 22: 835–841, 2011 doi:10.1093/annonc/mdq440 Published online 11 October 2010

Baseline nutritional evaluation in metastatic lung cancer patients: Mini Nutritional Assessment versus weight loss history



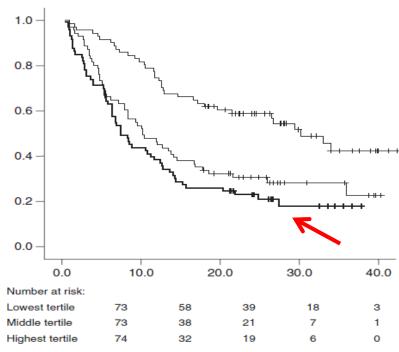


Figure 1 Survival curve representing survival duration in the patient cohort from time of diagnosis stratified according to tertiles of rate of weight loss. Thin line=lowest rate of weight-loss tertile with a median survival of 30.2 months; middle line = middle rate of weight-loss tertile with a median survival of 10.2 months; thick line = highest rate of weight-loss tertile with a median survival of 7.5 months (P < 0.0001, log-rank test).

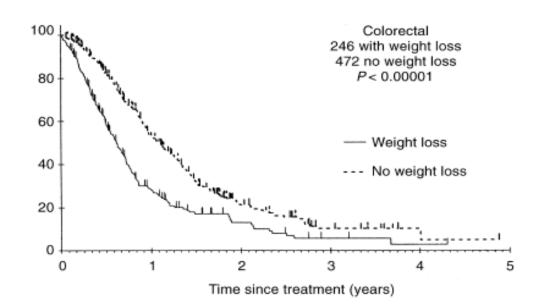
Ann Oncol. 2011, 22(4):835-41.

Weight change before presentation is associated with poorer outcomes in GI cancer



| | 51% weight | 49% weight | |
|--------------------|------------|-------------|----------|
| | stable | loss | |
| QoL score (0-100) | 76 | 59 | p<0.0001 |
| Stomatitis 1-4 | 39 % | 52 % | p<0.0001 |
| Treatment duration | 150 d | 120 d | p<0.0001 |
| Response rate | higher | lower | p=0.006 |
| Overall survival | 11.9 m | 7.6 m | p<0.0001 |

N=1555, (Age 18-84 y)

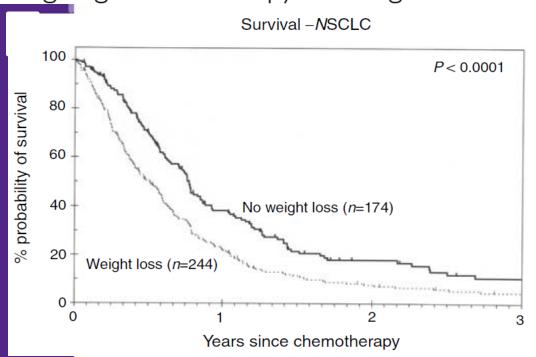




Weight loss is an independent PROGNOSTIC factor for survival in NSCLC



Do patients with weight loss have a worse outcome when undergoing chemotherapy for lung cancers?



Patients with weight loss and NSCLC (p=0.003) more frequently failed to complete at least three cycles of chemotherapy

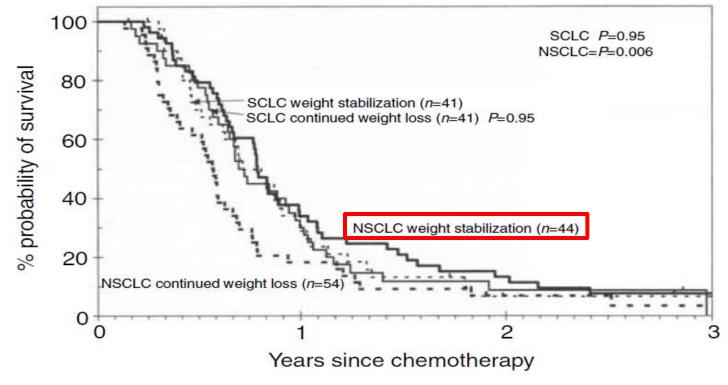
Prospectively collected data, stage III/IV NSCLC



Weight stabilization during chemotherapy contributes to higher survival



Do patients with weight loss have a worse outcome when undergoing chemotherapy for lung cancers?



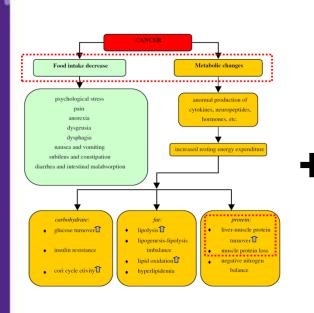
Prospectively collected data, stage III/IV NSCLC

British Journal of Cancer (2004) 90, 1905 – 1911 © 2004 Cancer Research UK All rights reserved 0007 – 0920/04 \$25.0



Standard chemotherapy & other cancer treatments reinforce cancer weight loss





Carmustine
Carboplatin
Cisplatin
5-Fluoruracil
Doxorubicin
Paclitaxel

Sorafinib Everolimus

Radiation Surgery



Weight loss at the start of treatment is associated with reduced response rates and increased toxicity and is included as one of the key Common Terminology criteria of Adverse events (CTCAE)

Prevalence of Side Effects of cancer treatments



| Treatment | Weight Loss | Fatigue | Nausea/ Vomiting | Oral Mucositis | Taste Alterations | Constipation |
|----------------------|----------------|--------------|---------------------|-------------------|----------------------|--------------|
| Overall % | 50%- 90% | 70%- 100% | 30%- 90% | 40%- 100% | 35%- 70% | 40%- 50% |
| Chemotherapy | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| Radiation | ✓ | ✓ | ✓ | ✓ | ✓ | |
| Surgery | ✓ | ✓ | ✓ | | | |
| Immunotherapy | ✓ | ✓ | | ✓ | | |



Cancer Cachexia



Definition and classification of cancer cachexia: an international consensus

THE LANCET Oncology

Kenneth Fearon*, Florian Strasser*, Stefan D Anker, Ingvar Bosaeus, Eduardo Bruera, Robin L Fainsinger, Aminah Jatoi, Charles Loprinzi, Neil MacDonald, Giovanni Mantovani, Mellar Davis, Maurizio Muscaritoli, Faith Ottery, Lukas Radbruch, Paula Ravasco, Declan Walsh, Andrew Wilcock, Stein Kaasa, Vickie E Baracos

Consensus findings Definition and diagnosis

Cancer cachexia is defined as a multifactorial syndrome characterised by an ongoing loss of skeletal muscle mass (with or without loss of fat mass) that cannot be fully reversed by conventional nutritional support and leads to progressive functional impairment. The pathophysiology is characterised by a negative protein and energy balance driven by a variable combination of reduced food intake and abnormal metabolism. Consensus statements for diagnosis are presented in the panel.

International consensus group: classification cancer cachexia in relation to outcomes

THE LANCET Oncology

| | Precachexia | Cachexia | Refractory cachexia |
|--------|---|---|---|
| Normal | | | Death |
| | Weight loss ≤5% Anorexia and metabolic change | Weight loss >5% or BM1<20 and weight loss >2% or sarcopenia and weight loss >2% Often reduced food intake/ systemic inflammation | Variable degree of cachexia Cancer disease both procatabolic and not responsive to anticancer treatment Low performance score <3 months expected survival |



Recommendation for early recognition of cachexia

Fearon KCH. Eur J Cancer, 2008 & Fearon et al, Lancet Oncol

Aopro et al, position paper of an ESO taskforce, 2014: Ann

Panel: Diagnosis of cancer cachexia

- Weight loss >5% over past 6 months (in absence of simple starvation); or
- · BMI < 20 and any degree of weight loss > 2%; or
- Appendicular skeletal muscle index consistent with sarcopenia (males <7.26 kg/m²; females <5.45 kg/m²)* and any degree of weight loss >2%†

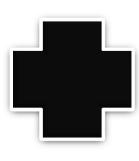
Cancer Cachexia



Assessment

Standard Parameters

Weight loss ≥ 5% over past 12 months (or BMI<20kg/m²)



THE LANCET Oncology

3/5 Parameters







A.S.P.E.N. Clinical Guidelines: Nutrition Support Therapy During Adult Anticancer Treatment and in Hematopoietic Cell Transplantation

ESPEN GUIDELINES

ESPEN Guidelines on Enteral Nutrition: Non-surgical oncology

Clinical Nutrition 28 (2009) 445-454



Contents lists available at ScienceDirect

Clinical Nutrition

journal homepage: http://www.elsevier.com/locate/clnu



Nutritional Interventions



Major goals of supportive nutrition

- Adjunctive to the specific oncology treatment goal
- maintain adequate nutritional status, body composition, performance status, immune function, and quality of life
- Stabilize or improve nutritional status as well as increasing the potential of a favorable response to therapy and enhancing recovery from any adverse effect of therapy
- early supportive nutritional intervention is to avoid irreversible nutritional and physiological deficits
- Weight loss in the cancer patient can often be prevented, but generally only of addressed proactively





ESPEN Guideline 2006- 2009; ASPEN Guideline 2009

Appropriate nutritional method

- Oral nutrition support (ONS)
- Tube feeding
- Parenteral nutrition (PN)

Adequate nutrition.

E: 25-30kcal/ kg/day

Protein: 1,2-1,5g/ kg/ day

(max 2g)

50% Energy not from protein

aspen____ Clinical Guidelines

A.S.P.E.N. Clinical Guidelines: Nutrition Support Therapy During Adult Anticancer Treatment and in Hematopoietic Cell Transplantation Supplements - Medicine

- Omega 3
- Drugs (steroids, progesterone, Cannabinoids, NSAIDS)

ESPEN GUIDELINES

ESPEN Guidelines on Enteral Nutrition: Non-surgical oncology

Nutrition in Cancer Care



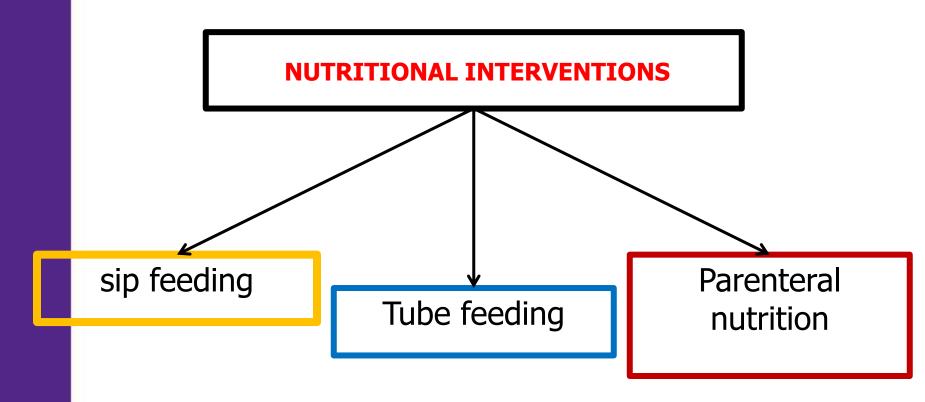
ESPEN Guideline 2006- 2009; ASPEN Guideline 2009

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|-----|------------------|
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| Cli | nical Guidelines |

ESPEN GUIDELINES

A.S.P.E.N. Clinical Guidelines: Nutrition Support Therapy During Adult Anticancer Treatment and in Hematopoietic Cell Transplantation

ESPEN Guidelines on Enteral Nutrition: Non-surgical oncology [☆]



Nutrition in Cancer Care



ESPEN Guideline 2006- 2009; ASPEN Guideline 2009

3.2. Is there an indication for EN during radiotherapy or combined radio-chemotherapy?

Yes. Use intensive dietary counselling and ONS to increase dietary intake (A) and to prevent therapy-associated weight loss and interruption of radiation therapy in patients undergoing radiotherapy of gastrointestinal or head and neck areas (A). If an obstructing head and neck or oesophageal cancer interferes with swallowing, EN should be delivered by tube (C). TF is also suggested if severe local mucositis is expected, which might interfere with swallowing, e.g. in intensive radiotherapy or in combined modality radio-chemotherapy regimens including radiation of throat or esophagus (C).



A.S.P.E.N. Clinical Guidelines: Nutrition Support Therapy During Adult Anticancer Treatment and in Hematopoietic Cell Transplantation



Chọn phương pháp dinh dưỡng



ESPEN Guideline 2006- 2009; ASPEN Guideline 2009

Clinical Nutrition 28 (2009) 445-454

Contents lists available at ScienceDirect

Clinical Nutrition

journal homepage: http://www.elsevier.com/locate/clnu



ESPEN Guidelines on Parenteral Nutrition: Non-surgical oncology

Nutritional support should be started if patient is undernourished or if it is anticipated that the patient will be unable to eat for more than seven days. It should also be started if an inadequate food intake (<60% of estimated energy expenditure) is anticipated for more than 10 days (Grade C). In such cases if nutritional support for any reason cannot be given through the enteral route, it has to be delivered by vein. A "supplemental" PN should substitute the difference between the actual oral/enteral intake and the estimated requirements (Grade C).

There is no rationale for giving PN if the nutrients intake by oral or enteral route is adequate, and for these reasons PN should not be administered in such conditions (Grade A).

Overview



- Weight loss to cancer-induced cachexia
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What can be Done??



- Conventional nutritional interventions have limited success
 - Standard oral nutritional products
 - Tube feeding
 - Total parenteral nutrition (TPN)

Conventional nutritional interventions do not address the underlying mechanism of Cancerinduced weight loss



What can be Done??



Clinical Nutrition 33 (2014) 749-753



Contents lists available at ScienceDirect

Clinical Nutrition





Randomized control trials

Randomized trial of the effects of individual nutritional counseling in cancer patients[☆]



Grith M. Poulsen a,c, Louise L. Pedersen a,c, Kell Østerlind b, Lene Bæksgaard b, Jens R. Andersen a,c,*

- ^a Department of Nutrition, Exercise and Sports, University of Copenhagen, DK-1958 Frederiksberg C, Denmark
- ^b Department of Oncology 5073, Rigshospitalet, DK-2100 Copenhagen Oe, Denmark
- ^c Nutrition Unit 5711, Rigshospitalet, DK-2100 Copenhagen Oe, Denmark

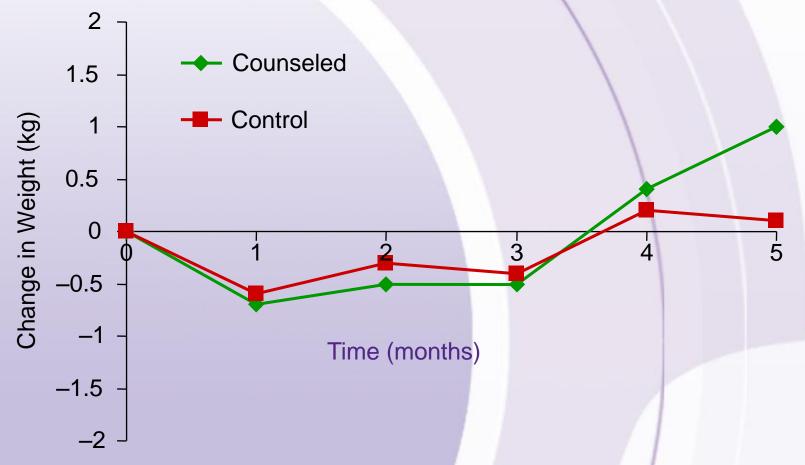
Conventional nutritional interventions *do not* address the underlying mechanism of Cancerinduced weight loss

Increasing Intake alone DOESN'T work





- •105 patients with small-cell lung, ovarian or breast cancer
- ·Significant increase in intake, but no significant weight gain



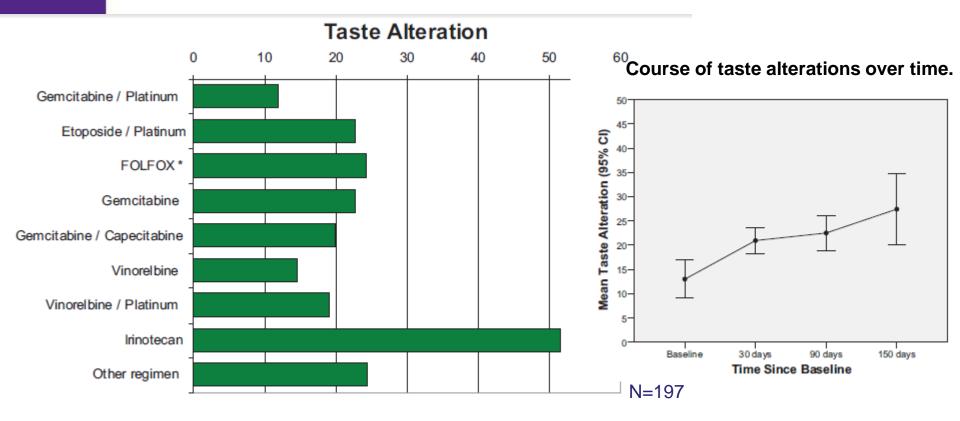
70% reported taste alterations (TA) during

& symptoms persist months after CT

VIỆT NAM HIM

Lung cancer 54%
Pancreatic cancer 19%
Colorectal cancer 26%
Age: 65 years; male 57%

included in study between day 0 and 30 of CT



Zabernigg et al, Taste alterations in in Cancer Patients receiving Chemotherapy, The Oncologist CME program 2010

Association between Taste alterations and QoL



EORTC QLQ – C30 + 2 additional questions

TAs are significantly associated with:

- Apetite loss
- Fatigue
- Nausea/vomiting
- Cognitive functioning



Challenges for intake



Taste alterations are common in cancer patients resulting from disease and/or treatment

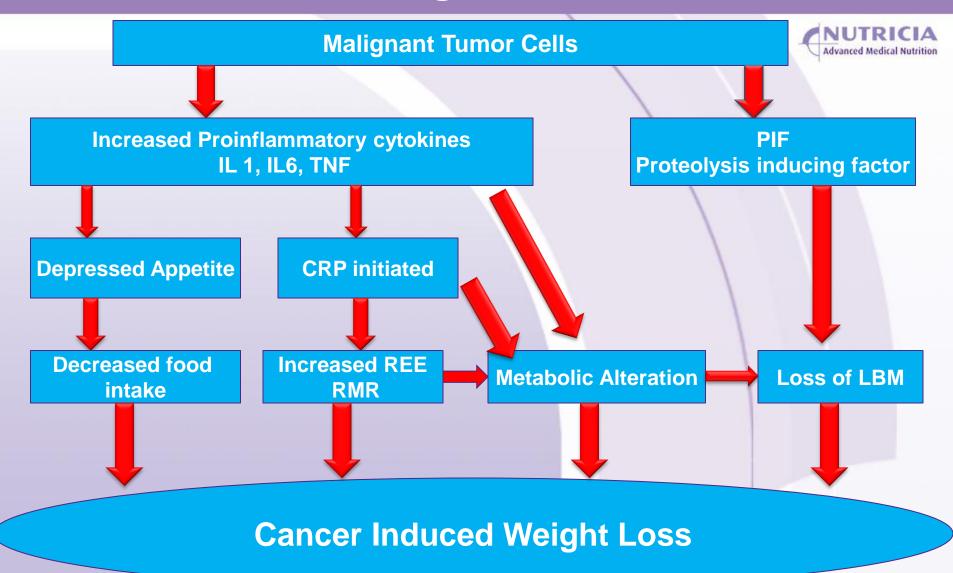
68% patients undergoing chemotherapy reported taste changes1:

- Food tastes like cardboard or sandpaper
- Too salty
- Too sweet
- Too sour
- Too bitter
- Metallic aftertaste

Prevalence of metallic taste ranged from 9.7 - 78% among patients with various cancers, chemotherapy treatments, and treatment phases

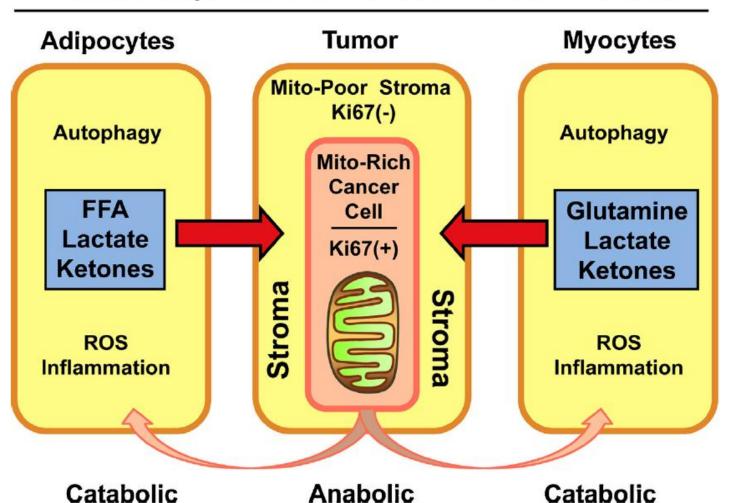
- ¹Wickham et al, 1999
- ² Ijpma et al, Cancer Treat Rev 2014

Mechanism of action Cancer induced weight loss



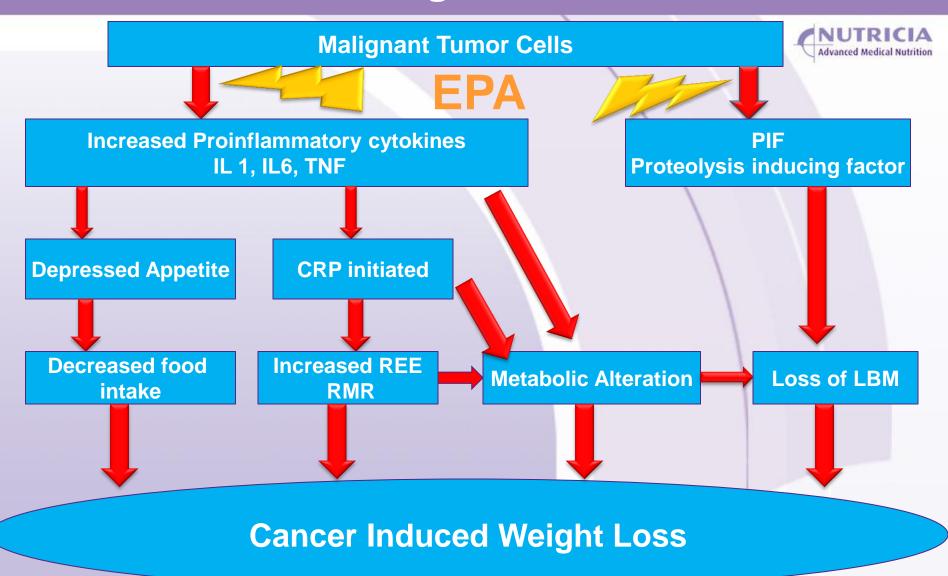
Catabolic cancer-associated fibroblasts transfer energy and biomass to anabolic cancer cells, fueling tumor growth

Multi-Compartment Metabolism in Cancer Cachexia



Martinez-Outschoorn UE et al. Sem Cancer Biol 2014; 25:47-60

Mechanism of action Cancer induced weight loss



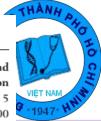


A.S.P.E.N. Clinical Guidelines: Nutrition Support Therapy During Adult Anticancer Treatment and in Hematopoietic Cell Transplantation

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Enteral Nutrition
Volume 33 Number 5
September/October 2009 472-500
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Parenteral and Enteral Nutrition
10.1177/0148607109341804

http://jpen.sagepub.com hosted at http://online.sagepub.com

8. ω-3 Fatty acid supplementation may help stabilize weight in cancer patients on oral diets experiencing progressive, unintentional weight loss. (Grade: B)



Clinical Study Model





Wigmore 1996
2 g EPA/day
Weight Stabilization

Barber 1999
Increase in weight and LBM

Wigmore 2000 6 g EPA/day Weight Stabilization

Barber 2001

Decrease in PIF and IL-6

aspen___ Clinical Guidelines

A.S.P.E.N. Clinical Guidelines: Nutrition Support Therapy During Adult Anticancer Treatment and in Hematopoietic Cell Transplantation Journal of Parenteral and Enteral Nutrition Volume 33 Number 5

tle or no increase in lean body mass. 102,105,106,111,112,116 A target dose of 2 g of eicosapentanoic acid daily appears appropriate. This may be administered as commercially available ω -3 enriched liquid nutritional supplements or as over-the-counter ω -3 fatty acid supplements (available in most pharmacies). Because these supplements are not commonly covered by health insurance, the cost of this intervention should be considered.



Improved body weight and performance after supplementation in newly diagnosed esophageal cancer patient Journal of Cachexia, Sarcopenia and Muscle 2015; 6: 32–44

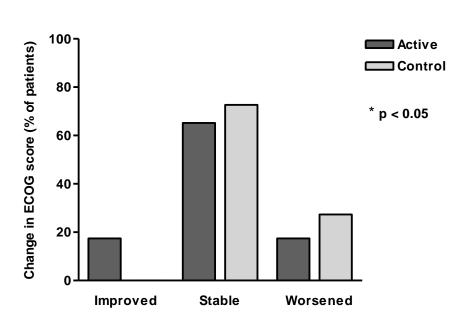
Body weight change

A * p < 0.05 1.75 1.50 (b) 1.25 1.00 1.00 0.75 0.25 -

Control

Active

Performance score



J Cachexia Sarcopenia Muscle. 2015, 6(1):32-44



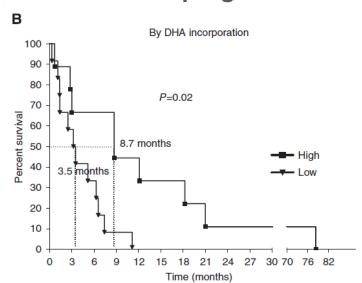
0.00

Improving outcome of chemotherapy of metastatic breast cancer by docosahexaenoic acid: a phase II trial

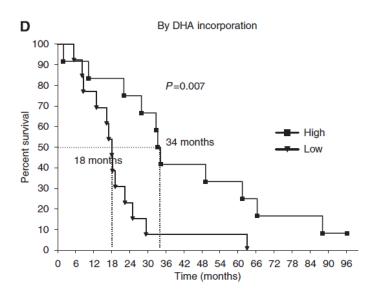


Improved outcome of FEC 75 chemotherapy in metastatic breast cancer

Time to progression



Overall survival





COLON

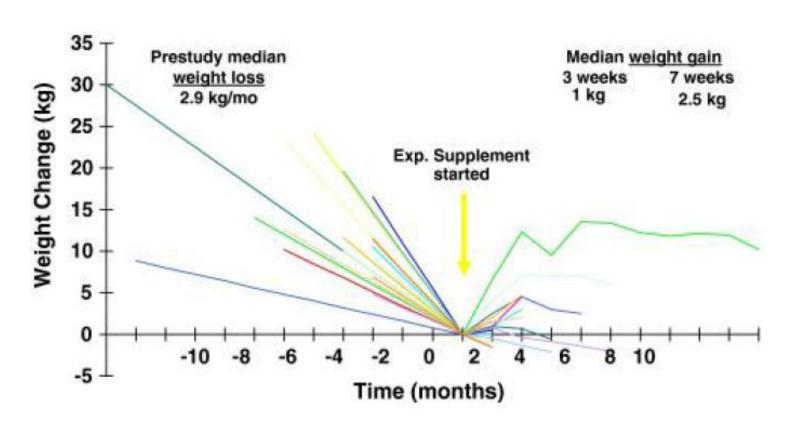
Effect of a protein and energy dense n-3 fatty acid enriched rsity of Medicine and Pharmacy oral supplement on loss of weight and lean tissue in cancer at Ho Chi Minh City, Vietnam



K C H Fearon, M F von Meyenfeldt, A G W Moses, R van Geenen, A Roy, D J Gouma, A Giacosa, A Van Gossum, J Bauer, M D Barber, N K Aaronson, A C Voss, M J Tisdale

Gut 2003;52:1479-1486

Cancer cachexia- ω3



Gut. 2003 52(10):1479-86.

ORIGINAL ARTICLE

Oral nutritional supplements containing n-3 polyunsaturated fatty acids affect quality of life and functional status in lung cancer patients during multimodality treatment: an RCT



BS van der Meij¹, JAE Langius¹, MD Spreeuwenberg², SM Slootmaker³, MA Paul⁴, EF Smit⁵ and PAM van Leeuwen⁴

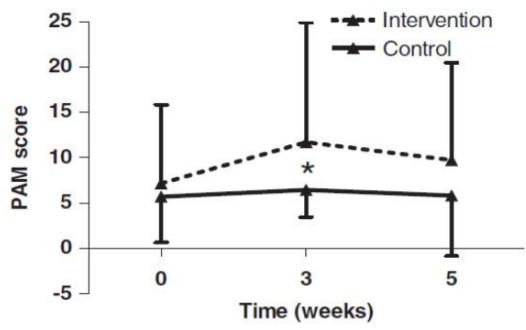


Figure 1. Physical activity (daily PAM score) over time for the I and C groups. Values are mean \pm s.d., baseline: n=12 (I), n=16 (C); week 3: n=13 (I) and n=17 (C); week5: n=8 (I), n=13 (C). *P < 0.05, difference between the I and C group (analysed by generalised estimating equations, with baseline value and sex as covariate).







Oral Nutritional Interventions in Malnourished Patients With Cancer: A Systematic Review and Meta-Analysis

Christine Baldwin, Ayelet Spiro, Roger Ahern, Peter W. Emery

Not reduce risk of death

| | Intervention | | No intervention | | Risk ratio | | Risk ratio |
|-------------------------------------|--------------|---------|-----------------|-------|------------|--------------------|--|
| Study or Subgroup | Events | Total | Events | Total | Weight | M-H, Fixed, 95% CI | M-H, Fixed, 95% CI |
| | | | | | | | |
| Baldwin et al. 2008a (17) | 29 | 90 | 9 | 32 | 8.0% | 1.15 [0.61-2.15] | - |
| Baldwin et al. 2008b (17) | 25 | 86 | 9 | 32 | 7.9% | 1.03 [0.54-1.97] | - |
| Baldwin et al. 2008c (17) | 22 | 86 | 9 | 32 | 7.9% | 0.91 [0.47-1.76] | |
| Elkort et al. 1980 (25) | 4 | 24 | 3 | 23 | 1.8% | 1.28 [0.32-5.10] | |
| Evans et al. 1987 (26) | 94 | 111 | 62 | 69 | 46.0% | 0.94 [0.84-1.05] | • |
| Isenring et al. 2004 (27) | 2 | 29 | 2 | 31 | 1.2% | 1.07 [0.16-7.10] | |
| Lovik et al. 1996 (28) | 3 | 28 | 0 | 24 | 0.3% | 6.03 [0.33-111.27] | |
| Moloney et al. 1983 (29) | 27 | 42 | 26 | 42 | 15.6% | 1.04 [0.75-1.44] | + |
| Ollenschlager et al. 1992(24) | 2 | 15 | 0 | 16 | 0.3% | 5.31 [0.28-102.38] | |
| Ovesen et al. 1993 (30) | 12 | 57 | 5 | 48 | 3.3% | 2.02 [0.77-5.33] | |
| Persson et al. 2002 (18) | 13 | 67 | 13 | 70 | 7.7% | 1.04 [0.52-2.09] | |
| Ravasco et al. 2005a (19) | 0 | 25 | 0 | 13 | | Not estimable | |
| Ravasco et al. 2005b (19) | 0 | 25 | 0 | 12 | | Not estimable | |
| Ravasco et al. 2005c (20) | 0 | 37 | 0 | 19 | | Not estimable | |
| Ravasco et al. 2005d (20) | 0 | 37 | 0 | 18 | | Not estimable | |
| Subtotal (95% CI) | | 759 | | 481 | 100.0% | 1.06 [0.92_1.22] | † |
| Total events | 233 | | 138 | | | | |
| Heterogeneity: $\chi^2 = 8.69$, df | r = 10 (P) | =.56);/ | $1^2 = 0\%$ | | | | |
| Test for overall effect: $Z = 0$. | 80 (P = .4) | 13) | | | | | |
| | | | | | | | |
| | | | | | | | 0.02 0.1 1 10 5 |
| nutritional inte | rven | tions | s and | | | | Favours no intervention Favours intervention |

Oral nutritional interventions and mortality meta-analysis.

J Natl Cancer Inst. 2012;104(5):371-85

University of Medicine and Pharmacy at Ho Chi Minh City, Vietnam



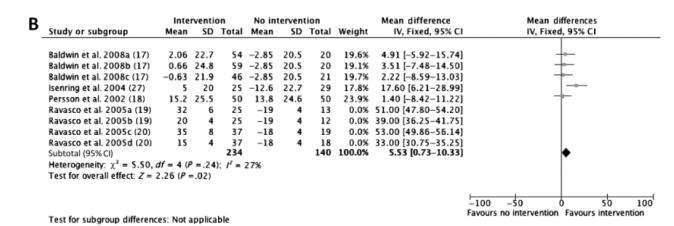
Oral Nutritional Interventions in Malnourished Patients With Cancer: A Systematic Review and Meta-Analysis

Christine Baldwin, Ayelet Spiro, Roger Ahern, Peter W. Emery

Improve the quality of life

Oral nutritional intervention and global quality of life metaanalysis

| Α | | Intervention | | No intervention | | | Mean difference | | Mean difference | |
|----|----------------------------------|----------------|--------|-----------------|------------|--------|-----------------|--------|---------------------|--|
| ٦. | Study or subgroup | Mean | SD | Total | Mean | SD | Total | Weight | IV, Random, 95% CI | IV, Random, 95% CI |
| | | | | | | | | | | |
| | Baldwin et al. 2008a (17) | 2.06 | 22.7 | 54 | -2.85 | 20.5 | 20 | 10.5% | 4.91 [-5.92-15.74] | → |
| | Baldwin et al. 2008b (17) | 0.66 | 24.8 | 59 | -2.85 | 20.5 | 20 | 10.4% | 3.51 [-7.48-14.50] | - |
| | Baldwin et al. 2008c (17) | -0.63 | 21.9 | 46 | -2.85 | 20.5 | 21 | 10.5% | 2.22 [-8.59-13.03] | + |
| | Isenring et al. 2004 (27) | 5 | 20 | 25 | -12.6 | 22.7 | 29 | 10.3% | 17.60 [6.21-28.99] | |
| | Persson et al. 2002 (18) | 15.2 | 25.5 | 50 | 13.8 | 24.6 | 50 | 10.7% | 1.40 [-8.42-11.22] | 1 |
| | Ravasco et al. 2005a (19) | 32 | 6 | 25 | -19 | 4 | 13 | 11.9% | 51.00 [47.80-54.20] | * |
| | Ravasco et al. 2005b (19) | 20 | 4 | 25 | -19 | 4 | 12 | 11.9% | 39.00 [36.25-41.75] | + |
| | Ravasco et al. 2005c (20) | 35 | 8 | 37 | -18 | 4 | 19 | 11.9% | 53.00 [49.86-56.14] | * |
| | Ravasco et al. 2005d (20) | 15 | 4 | 37 | -18 | 4 | 18 | 12.0% | 33.00 [30.75-35.25] | |
| | Subtotal (95% CI) | | | 358 | | | 202 | 100.0% | 24.02 [14.33-33.72] | • |
| | Heterogeneity: $\tau^2 = 203.42$ | $: \chi^2 = 3$ | 49.77 | df = i | 8 (P < .0) | 0001); | $I^2 = 98$ | 3% | | |
| | Test for overall effect: $Z = 4$ | .86 (P | <.0000 | 1) | | | | | | |
| | | | | | | | | | | |
| | | | | | | | | | | -100 -50 0 50 100 |
| | | | | | | | | | | Favours no intervention Favours intervention |



J Natl Cancer Inst. 2012;104(5):371-85

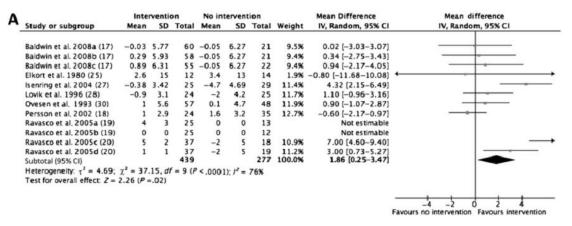


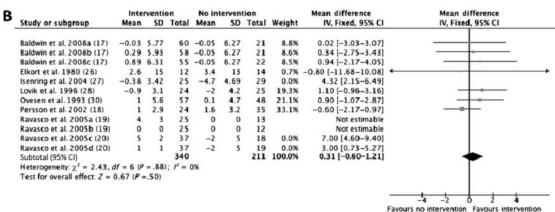
Oral Nutritional Interventions in Malnourished Patients With Cancer: A Systematic Review and Meta-Analysis

Christine Baldwin, Ayelet Spiro, Roger Ahern, Peter W. Emery

Weight gain

Oral nutritional interventions and weight gain meta-analysis.





J Natl Cancer Inst. 2012;104(5):371-85

University of Medicine and Pharmacy at Ho Chi Minh City, Vietnam

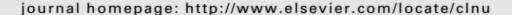


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Original article

Muscle protein synthesis in cancer patients can be stimulated with a specially formulated medical food*

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Development of Forticare: Adapted Medical Nutrition for cancer patients

FortiCare is nutritionally complete Provides high protein and is EPA/DHA enriched

- EPA to attenuate cachexia, reduce inflammation and support immune function
- Energy and protein to meet increased needs of cancer patients
- Adapted taste, small volume (for improved compliance)





Forticare Clinical trial in Vietnam



Clinical trial: Randomized controlled trial (RTC) on 60 patients

CILW: 3,3 kg per month. Colorectal cancer patients with malnutrition or cachexia.

Hospitals:

Department of Surgery C – K Hospital. Clinical Nutrition Center of Bach Mai Hospital Oncology and Palliative Care Unit, Hanoi Medical University Hospital

Date: $12/2012 \rightarrow 6/2015$



Forticare Clinical trial in Vietnam



Nutrition intervention with EPA (2g / day): average weight gain of 3 kg/patient after 8 weeks of treatment.

Nutrition interventions EPA contributes to improve the quality of life for patients:

100% appetite (Delicious and good taste)

Weight gain: 3kg (8weeks)

Increase the size arm circumference

Improves albumin/blood



Forticare Clinical trial in Vietnam



EPA treatment for colorectal cancer patients:

Nutrition support should be continuous treatment.

Protein: 1,5 - 1,7g / kg / day

Energy: 35 kcal / kg / day

EPA 2g/day

Recommendation: The EPA should be included in the treatment for cancer patients to prevent weight loss and cachexia.

ESPEN Guidelines on Enteral Nutrition: Non-surgical oncology



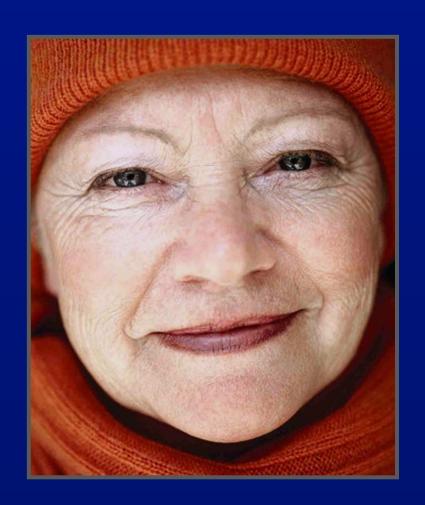
J. Arends^{a,*}, G. Bodoky^b, F. Bozzetti^c, K. Fearon^d, M. Muscaritoli^e, G. Selga^f, M.A.E. van Bokhorst-de van der Schueren^g, M. von Meyenfeldt^h, DGEM: A G. Zürcher, R. Fietkau, E. Aulbert, B. Frick, M. Holm, M. Kneba, H.J. Mestrom, A. Zander

Clinical Nutrition (2006) **25**, 245–259

The efficacy of treatment with EPA/ONS appears to be critically dependent on the patients' compliance. In addition to anorexia, patients' compliance with prescribed high-energy and high-protein EPA/ONS is limited by the frequently complained unpleasant aftertaste. Therefore, it will be necessary to improve the palatability of EPA/ONS in order to improve patients' compliance with treatment and hopefully its effectiveness.

Any food that is not consumed is never nutritious!

Prof Jeya Henri





WAYS TO PROVIDE > 2 GRAMS OF EPA / DAY

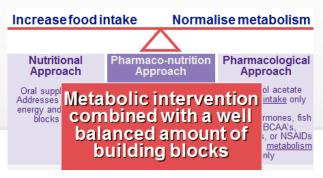
Eat large amounts of fatty fish

 Herring, Salmon, Tuna, Mackerel, Sardines....

Fish oil capsules

Emulsified oils combined with macro- and micro- nutrients





FORTICARE: A CONVENIENT WAY TO PROVIDE THE ESSENTIAL NUTRITION FOR CANCER PATIENTS

Oncology Adapted Medical Nutrition:

- ✓ Energy dense
- ✓ High in protein
- ✓ Nutritionally complete
- ✓ Low GI
- ✓ Small volume
- ✓ Low viscosity
- ✓ Enriched with EPA
- ✓ Excellent taste

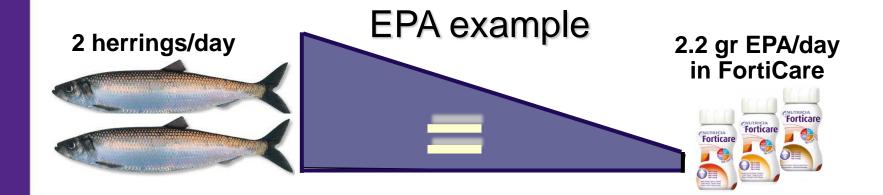


FULFILL PATIENTS NEEDS IN A CONVENIENT WAY

High energy High amount of protein High quality of protein All other macro-nutrients Dietary fibers Balanced micro-nutrients

EPA (6 ounce of fat fish)

| Convenience |
|---------------|
| Small volume |
| Complete |
| Balanced |
| Low viscosity |
| Tasty |





Oncology Adapted Medical Nutrition





WHY:

 To improve the outcome of your cancer treatment and QOL for the patient.

WHEN:

Implement screening for nutritional status of every newly diagnosed cancer patient and consider intervention options in every stage of the disease.

HOW:

Intensive nutritional support with nutrients and metabolic modulators in a convenient and palatable way.

Comparison





FortiCare

FortiCare





Overview



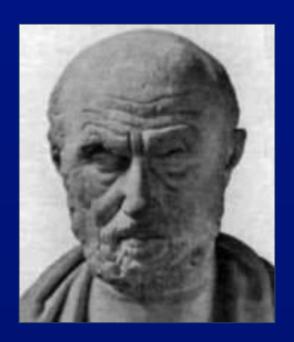
- Weight loss to cancer-induced cachexia
- Specific nutrition for cancer patients -Nutrition in Cancer Care
- Summary

Summary



- Early detection and signs of cancer-induced weight loss in every stage of the disease.
- Cancer cachexia affects clinical outcome, increased doselimiting toxicities, receive less treatment (dosage), and treatment interruption.
- Conventional nutritional interventions do not address the underlying mechanism of cancer-induced weight loss and cancer cachexia
- High energy, High protein and high EPA (2g/day) are recommended for cancer patents.
- Considered taste alterations, flavor and the patients' compliance with treatment.
- Hyperglycemia is common in cancer patients and low Glycemic index is suitable for cancer patients.

"Let medicine be thy food, and food be thy medicine."



Hippocrates of Cos, Greece 460-377 B.C.

CẢM ƠN SỰ CHÚ Ý LẮNG NGHE CỦA QUÝ VỊ!

Thank You

For Your Attention!!!